

PARTICULARS TO APPEAR ON THE OUTER PACKAGE

10 x 2 ml dose Folding Carton

10 x 2 ml dose Booklet Label

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Equip T suspension for injection

2. STATEMENT OF ACTIVE SUBSTANCES

Each 2 ml dose contains:

Concentrated immunopurified tetanus toxoid (≥ 30 IU/ml) adsorbed onto aluminium phosphate.

3. PACKAGE SIZE

10 x 2 ml

4. TARGET SPECIES

Horses.

5. INDICATIONS

6. ROUTES OF ADMINISTRATION

Intramuscular use.

7. WITHDRAWAL PERIODS

Zero days.

8. EXPIRY DATE

Exp. {mm/yyyy}

9. SPECIAL STORAGE PRECAUTIONS

Store in a refrigerator.
Protect from light.
Do not freeze.
Keep the container in the outer carton.

10. THE WORDS “READ THE PACKAGE LEAFLET BEFORE USE”

Read the package leaflet before use.

11. THE WORDS “FOR ANIMAL TREATMENT ONLY”

For animal treatment only.

12. THE WORDS “KEEP OUT OF THE SIGHT AND REACH OF CHILDREN”

Keep out of the sight and reach of children.

13. NAME OF THE MARKETING AUTHORISATION HOLDER

Zoetis UK Limited

14. MARKETING AUTHORISATION NUMBERS

Vm 42058/5232

15. BATCH NUMBER

Lot {number}

MINIMUM PARTICULARS TO APPEAR ON SMALL IMMEDIATE PACKAGING
UNITS
2 ml Vial Label 2 ml Syringe Label

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Equip T

2. QUANTITATIVE PARTICULARS OF THE ACTIVE SUBSTANCES

Each 2 ml dose contains:

C.tetani toxiod

3. BATCH NUMBER

Lot {number}

4. EXPIRY DATE

Exp. {mm/yyyy}

PARTICULARS TO APPEAR ON THE PACKAGE LEAFLET:

PACKAGE LEAFLET

1. Name of the veterinary medicinal product

Equip T suspension for injection

2. Composition

Each 2 ml dose contains:

Active substance:

Immunopurified Tetanus Toxoid ≥ 30 IU/ml*

* IU: International units

Clear liquid suspension above a whitish/grey sediment which resuspends readily on shaking.

3. Target species

Horses.

4. Indications for use

For the active immunisation of horses of 5 months of age or older against tetanus to prevent mortality.

Onset of immunity: within 2 weeks of completion of the primary course

Duration of immunity: 3 years.

5. Contraindications

None.

6. Special warnings

Vaccinate healthy animals only.

Special precautions for safe use in the target species:

The efficacy of active immunisation of young foals against equine influenza will be influenced by the level of maternally derived antibodies. This will vary between individuals due to a number of factors, e.g. the immune status of the dam; adequacy of colostral intake by the foal, etc. The vaccine should not be used in foals below 5 months of age, and foals should not be vaccinated until maternally derived antibodies have fallen below protective levels.

In any animal population, there may be a small number of individuals which fail to respond fully to vaccination. Successful vaccination depends upon correct storage and administration of the vaccine and the ability of the animal to respond. This can be influenced by such factors as genetic constitution, intercurrent infection, age, nutritional status, concurrent drug therapy and stress.

Special precautions to be taken by the person administering the veterinary medicinal product to animals:

The product should be administered by respecting appropriate (aseptic) injection technique.

In case of accidental self-injection, seek medical advice immediately and show the package leaflet or the label to the physician.

Pregnancy and lactation:

The vaccine may be used in pregnant mares which have been vaccinated against tetanus before pregnancy.

Heavily pregnant mares should not be subject to undue stress when vaccinated.

Interaction with other medicinal products and other forms of interaction:

No information is available on the safety and efficacy of this vaccine when used with any other veterinary medicinal product. A decision to use this vaccine before or after any other veterinary medicinal product therefore needs to be made on a case by case basis.

Overdose:

Accidental overdosage is unlikely to cause any reactions other than those described in section 7.

Major incompatibilities:

Do not mix with any other veterinary medicinal product.

7. Adverse events

Horses.

Rare (1 to 10 animals / 10,000 animals treated):
Injection site swelling ^{1,3} Stiffness ¹ Elevated temperature ^{1,2}
Very rare (<1 animal / 10,000 animals treated, including isolated reports):
Injection site pain Hypersensitivity reaction ⁴

¹ This condition normally resolves by the day following vaccination.

² Mild, transient, typically 9-12 hours post vaccination.

³ Local, small (10-20 mm in diameter), soft, non-painful.

⁴ In the event of an allergic or anaphylactic reaction, immediate treatment should be given with a soluble glucocorticoid intravenously or adrenalin intramuscularly.

Reporting adverse events is important. It allows continuous safety monitoring of a veterinary medicinal product. If you notice any side effects, even those not already listed in this package leaflet, or you think that the medicine has not worked, please contact, in the first instance, your veterinarian. You can also report any adverse events to the marketing authorisation holder using the contact details at the end of this leaflet, or via your national reporting system at:

Website: <https://www.gov.uk/report-veterinary-medicine-problem/animal-reacts-medicine>
e-mail: adverse.events@vmd.gov.uk

8. Dosage for each species, routes and method of administration

Dose: 2 ml.

Administration: Equip T should be shaken thoroughly before use and administered by deep intramuscular injection.

9. Advice on correct administration

Primary vaccination

Two injections of 2 ml with an interval of 4-6 weeks between them

Booster vaccination

One dose 36 months after the primary course, repeated at intervals of up to 36 months.

10. Withdrawal periods

Zero days.

11. Special storage precautions

Keep out of the sight and reach of children.

Store in a refrigerator (2 °C - 8 °C).

Protect from light.

Do not freeze.

Keep the container in the outer carton.

Do not use this veterinary medicinal product after the expiry date which is stated on the label. The expiry refers to the last day of that month.

12. Special precautions for disposal

Medicines should not be disposed of via wastewater or household waste.

Use take-back schemes for the disposal of unused veterinary medicinal product or waste materials derived thereof in accordance with local requirements and with any applicable national collection systems. These measures should help to protect the environment.

Ask your veterinary surgeon how to dispose of medicines no longer required.

13. Classification of veterinary medicinal products

Veterinary medicinal product subject to prescription.

14. MARKETING AUTHORISATION NUMBERS AND PACK SIZES

42058/5232

Type I glass vial with chlorobutyl rubber stopper and aluminium overseal.
Packaging: Box of 10 single-dose vials. Each box contains 10 sterile disposable 2 ml syringes and 10 sterile needles.

Type I glass syringe closed with bromobutyl rubber plunger stopper and tip cap.
Packaging: Box of 10 single-dose prefilled syringes with needles

Not all pack sizes may be marketed.

15. PID link (Do not print heading)

[The following statement must be included where reference to the European Union Product Database is included on the product information. This statement is relevant to both UK(GB) and UK(NI) products:]

Find more product information by searching for the 'Product Information Database' on www.gov.uk.

16. Contact details

Marketing authorisation holder and contact details to report suspected adverse reactions:

Zoetis UK Limited
1st Floor, Birchwood Building
Springfield Drive
Leatherhead
Surrey
KT22 7LP
Phone: +44 (0) 345 300 8034

Manufacturer responsible for batch release:

Zoetis Belgium
Rue Laid Burniat 1
1348 Louvain-La-Neuve
Belgium

17. Other information

Equip T stimulates active immunity against tetanus.

POM-V

Gavin Hall
Approved: 11 December 2025